

GenCore version 4.5
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SUMMARIES

12	282	95.9	618	18	ANX33945	Human c-IAP1. Hom
13	282	95.9	618	18	ANW15583	Human cellular lnh
14	270	91.8	618	18	ANW15583	Human apoptosis in
15	270	91.8	618	19	ANW69396	Human IAP-2 prote
16	269	91.5	600	19	ANW69398	Murine IAP-1 proce
17	269	91.5	612	18	ANW15555	Murine c-IAP. Mur
18	269	91.5	612	19	ANW62939	Murine IAP-2 prot
19	263	89.5	591	18	ANW15586	Mouse apoptosis in
20	259	88.1	602	18	ANW15587	Mouse apoptosis in
21	210	85.6	210	22	ANM25285	Human protein in
22	193	65.6	280	22	ANB3178	Amino acid sequen
23	193	65.6	298	21	ANY61807	A human proliferat
24	193	65.6	298	21	ANY6182	Human inhibitor of
25	177	60.2	496	18	ANW19145	Mouse inhibitor of
26	177	60.2	497	18	ANW19581	Human apoptosis in
27	177	60.2	497	19	ANW69394	Human XIAP protein
28	177	60.2	497	21	ANY99895	Human X-I-linked inh
29	177	60.2	497	21	ANY95451	Human XIAP protein
30	173	58.8	236	21	AYX81440	Human TIP (an inh
31	173	58.8	236	22	ANB00365	Human IAP-like pro
32	173	58.8	236	22	ANB00366	Chimpanzee IAP-like
33	173	58.8	1232	17	ANH98216	Neuronal apoptosis
34	173	58.8	1295	20	AYA10480	Gonadotropic hormo
35	173	58.8	1295	20	AYX0940	Human apoptosis in
36	173	58.8	1403	18	ANW20032	Neuronal apoptosis
37	173	58.8	1403	18	ANW20033	Neuronal apoptosis
38	173	58.8	1403	20	AYV14079	Gonadotropic hormo
39	173	58.8	1403	20	AYX05359	Human apoptosis in
40	173	58.8	1403	21	AYA89053	Human MAP protein
41	172	58.5	236	22	ANR00167	Gorilla IAP-like p
42	103	55.4	434	22	ANB48195	Drosophila mutant
43	103	55.4	438	22	ANB48188	Drosophila wild-type
44	163	55.4	438	22	ANB48189	Drosophila mutant
45	163	55.4	438	22	ANB48190	Drosophila mutant

PS Claim 3: Page 25; 35PP; English.

XX CC The human cellular inhibitor of apoptosis proteins (c-IAP1/2 - AAT61590/61591) comprise a series of defined structural domain repeats and/or a RING finger domain; in particular, at least two of CC a first domain repeat (AAW13547 or AAW13548), a second domain repeat (AAW13549 or AAW13550), and a third domain repeat (AAW13551 or AAW13552) and/or a RING finger domain (AAW13553 or AAW13554), or a consensus sequences derived from these human genes.

CC They can also be used in conditions requiring a reduction in CC apoptosis.

CC They can also be used in conditions requiring a reduction in CC apoptosis.

XX Sequence 48 AA;

SQ

Query Match	Score	Length
Best Local Similarity	100.0%	48
Matches	48; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
Oy	1 PEQLASAGFVYVGNSDDVKKCFCGDLRKWEGDDPWQHAKWKPRLCE 48	
Db	1 peqlasagfvyvgnsddvkkcfcgdlrkwesggdpwqhwkprlce 48	

RESULT 2

ID AAW1947 standard; Protein; 604 AA.

XX AC AAW1947;

XX DT 16-SEP-1997 (first entry)

DE Human inhibitor of apoptosis protein homologue MIHC.

XX KW Inhibitor of apoptosis protein; IAP; mammalian IAP homologue; MIHC;

KW degenerative disease; infectious disease; autoimmune disease;

KW cancer; therapy; diagnosis.

XX OS Homo sapiens.

XX FH Location/Qualifiers

FT Key 29..97

FT Region /label= BIR

FT Region 169..236

FT Region /label= BIR

FT Region 255..323

FT Region /label= BIR

FT Region 556..593

FT Region /label= RING_finger

PN W09723501-A1.

XX PD 03-JUL-1997.

XX PR 20-DEC-1996; 96WO-AU00827.

PR 22-DEC-1995; 95AU-007275.

XX PA (AMPA-) AMRAD OPERATIONS PTY LTD.

XX PI VAUX DL;

XX DR WPI; 1997-350966/32.

DR N-PSDB; AAT72712.

XX PT Isolated protein homologues of viral inhibitors of apoptosis - used to modulate apoptosis for treatment of degenerative, Infectious or

PT autoimmune diseases and cancer

XX PS Claim 9; Page 58-62; 136PP; English.

XX CC Mammalian IAP homologue C (MIHC) (AAW1947) is a human homologue of baculovirus inhibitor of apoptosis protein (IAP). Its amino acid sequence was deduced from a cDNA clone (see also AAT7212) isolated from a human foetal liver cDNA library using primers based on human EST sequences that resembled the BIR repeats of orygia pseudotisgrata polydendrosis virus IAP. IAP homologues (see also AAW19745..46 and AAW19748..52) and their derivatives and chemical analogues can be used in methods for modulating apoptosis in animal cells, specifically for treatment, by inhibition, of degenerative and infectious disease or, by promotion, of cancer and autoimmune disease.

XX SQ Sequence 604 AA;

Query Match Similarity 100.0%; Score 294; DB 18; Length 604;

Best Local Similarity 100.0%; Pred. No. 2; 7e-28; Matches 48; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 PEQLASAGFVYVGNSDDVKKCFCGDLRKWEGDDPWQHAKWKPRLCE 48

Db 273 peqlasagfvyvgnsddvkkcfcgdlrkwesggdpwqhwkprlce 320

RESULT 3

ID AAW19582

XX AC AAW19582;

XX DT 02-SEP-1997 (first entry)

XX DE Human apoptosis inhibitor HIAP-1.

XX KW Apoptosis inhibitor; HIAP-1; HIV; AIDS; neurodegeneration; KW myelodysplastic syndrome; ischaemia; myocardial infarction; stroke; KW reperfusion injury; toxin-induced liver disease; gene therapy; KW diagnosis.

XX OS Homo sapiens.

XX FH Location/Qualifiers

FT Key 29..96

FT Domain /label= BIR-1

FT Domain 169..235

FT Domain /label= BIR-2

FT Domain 255..322

FT Domain /label= BIR-3

FT Domain 546..591

FT Domain /label= Ring_zinc_finger

PN W09706255-A2.

XX PD 20-FEB-1997.

XX PR 05-AUG-1996; 96WO-1B01022.

XX PR 22-DEC-1995; 95US5-0576956.

PR 04-AUG-1995; 95US-0511485.

XX PA (UYOT-) UNIV OTTAWA.

XX PI Baird S, Korneluk RG, Liston P, Mackenzie AE;

XX DR WPI; 1997-142462/14.

DR N-PSDB; AAT70837.

XX PT Nucleic acid encoding an inhibitor of apoptosis polypeptide - used to inhibit apoptosis in e.g. HIV or AIDS patients, and for detection

PT of susceptibility to apoptotic disease
 XX
 PS Claim 27; Page 72-74; 219PP; English.

XX Human XIAP, HIAP-1 and HIAP-2 and murine M-XIAP, M-HIAP-1 and
 CC M-HIAP-2 (AAW1981-86) are a new class of mammalian proteins that
 CC are inhibitors of apoptosis (IAP) and which are characterized by
 CC the presence of a ring finger domain (see also AAW1587) and at
 least one BIR (baculovirus IAP repeat) domain (see also AAW19568).
 The IAP amino acid sequences were deduced from cDNA clones (AAW7033/7
 CC and AAW7038) from a human liver library. IAP polypeptides can be
 expressed in host cells (in vitro or in vivo) and used in methods
 CC for treating diseases and disorders involving apoptosis, esp. in a
 CC human diagnosed as HIV-positive or as having AIDS, aitis, a
 CC ischemic injury, selected from myocardial infarction, stroke,
 CC neurodegenerative diseases a myelodysplastic syndrome or an
 CC reperfusion injury, or a toxin-induced liver disease.
 XX Sequence 604 AA:

Query Match Similarity 100.0%; Score 294; DB 18; Length 604;
 Best Local Similarity 100.0%; Pred. No. 2, 7e-28;
 Matches 48; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 273 pegasasgffyvvqnsdvdvcfcfdggirclrwesgdpwvhakwfprce 320

RESULT 4
 AAW1346 standard; Protein; 604 AA.
 ID AAW1346; AC AAW13546; DT 22-JUL-1997 (first entry)
 DE Human c-IAP2.
 XX
 KW IAP; inhibitor; apoptosis; RING finger domain; restinosis;
 KW myocardial infarction; nephritis; HIV.
 XX Homo sapiens.
 OS XX
 PN WO9706182-A1.
 PD 20-FEB-1997.
 PR 06-AUG-1996; 96WO-US12860.
 PR 08-DEC-1995; 95US-0559749.
 PR 08-AUG-1995; 95US-0552946.
 PA (TULA-) TULARIK INC.
 PI goedel DV, Rothe M;
 XX DR WPI; 1997-154209-14.
 DR N-PSBB; AAV5039.

PT Nucleic acids encoding cellular inhibitor of apoptosis proteins - useful for apoptosis regulation in cells to reduce or increase
 PT apoptosis and for pharmacological screening
 XX Disclosure, Page 21-23; 35pp; English.

XX The human cellular inhibitor of apoptosis proteins (c-IAP1/2 -
 CC AAT61590/T61591) comprise a series of defined structural domain
 CC repeats and/or a RING finger domain, in particular, at least two of
 CC a first domain repeat (AAW13547 or AAW13548), a second domain repeat
 CC (AAW13549 or AAW13550), and a third domain repeat (AAW13551 or AAW13552)
 CC and/or a RING finger domain (AAW13553 or AAW13554), or a consensus

CC sequences derived from these human genes.
 CC The nucleic acid is used for recombinant produc. of human cellular
 CC inhibitor of apoptosis protein which modulates apoptosis
 CC regulation. The nucleic acids are useful in therapies where
 CC increased cell-specific apoptosis is desired e.g. in retinosis,
 CC inflammatory disease states, myocardial infarction, glomerular
 CC nephritis, transplant rejection and infectious diseases, e.g. HIV.
 CC They can also be used in conditions requiring a reduction in
 CC apoptosis.

XX Sequence 604 AA:

Query Match Similarity 100.0%; Score 294; DB 18; Length 604;
 Best Local Similarity 100.0%; Pred. No. 2, 7e-28;
 Matches 48; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 273 pegasasgffyvvqnsdvdvcfcfdggirclrwesgdpwvhakwfprce 320

RESULT 5
 AAW69295 standard; Protein; 604 AA.
 ID AAW69295; AC AAW69295; DT 13-NOV-1998 (first entry)
 DE Human HIAP-1 protein.
 XX Inhibitor of apoptosis protein; apoptosis enhancer; IAP polypeptide;
 KW proliferative disease; IAP; therapy; cancer; HIAP-1 protein.
 XX OS Homo sapiens.
 PN WO9835693-A2.
 PD 20-AUG-1998.
 PR 13-FEB-1998; 98WO-1B00781.
 PR 13-FEB-1997; 97US-080929.
 PR (UYON-) UNIV OTTAWA.
 XX
 PI Baird S, Korneluk R, Liston P, Mackenzie AE, Pratt C;
 PI Tsang B;
 XX DR WPI; 1998-467164/40.
 DR N-PSBB; AAV5039.

PT Inducing apoptosis in proliferative mammalian cells with inhibitor
 PT of IAP or NAIP polypeptide - also methods for prognosis based on
 PT presence of IAP and NAIP, specifically applied to cancers involving
 PT p53 mutations
 XX Disclosure, Fig 2; 147PP; English.

XX This sequence is the human HIAP-1 protein, which is a inhibitor of
 CC apoptosis protein (IAP), and can be used in the method of the invention.
 CC The method is for enhancing apoptosis in cells from a mammal with
 CC proliferative disease by treatment with a compound that inhibits
 CC biological activity of an IAP or NAIP polypeptide. The inhibitor
 CC compounds are used to treat proliferative diseases, specially cancers of
 CC ovary, breast, pancreas, lymph nodes, skin, blood, lung, brain, kidney,
 CC liver, esophagus, thyroid, central nervous system, prostate, colon,
 CC rectum, cervix or endometrium, particularly to increase their sensitivity
 CC to chemotherapeutic agents. High levels of the IAP or NAIP proteins are
 CC detected in many cancers and are associated with poor prognosis,
 CC resistance to chemotherapeutic agents and mutations in p53 (it is
 CC suggested that wild-type p53 suppresses transcription of the IAP or NAIP

CC genes). Transgenic animals are used for testing the effects of antisense oligonucleotides and for screening for the inhibitors.

CC
XX
SQ Sequence 604 AA;

Query Match Similarity 100.0%; Score 294; DB 19; Length 604;
Best Local Similarity 100.0%; Pred. No. 2.7e-28; Mismatches 0; Indels 0; Gaps 0;
Matches 48; Conservative 0;

Qy 1 PEGIASGFYVGNSDVKCFCGGLRCMFGSDPPWQHAKWPRCE 48
Db 273 pegasagfyyvgnsdavkfcfcodgglcweesgdapwqhakwprce 320 .

RESULT 6

AAY52703
ID AAY52703 standard; Protein: 604 AA.
XX
AC XX
XX
DT 26-JAN-2000 (first entry)

DE Human cellular inhibitor of apoptosis-2 protein.
KW identification; genetic target; gene modulation; human;
antisense oligonucleotide; phosphorothioate; target validation;
nucleotide sequence-based technology; antisense drug discovery;
OS Homo sapiens.
XX
PN WO9953101-A1.
XX
PD 21-OCT-1999.
XX
PR 13-APR-1999; 99WO-US08268.
PR 13-APR-1998; 98US-0061493.
PR 28-APR-1998; 98US-006763B.
XX
(ISIS-) ISIS PHARM INC.
PI Cowser LM, Baker BF, McNeill J, Freier SM, Sasnor HM, Brooks DG;
XX
PI Ohasi C, Wyatt JR, Borchers AH, Vickers TA;
XX
DR WPI; 1999-620446/53.
DR N-PSDB; AA241005.
XX
PT Identifying compounds which modulate expression of nucleic acids, used
PT to provide compounds having defined physical, chemical or bioactive
PT properties, e.g., antisense activity -
PS Example 20; Page 197-202; 26pp; English.

A method has been developed or defining a set of compounds that modulate the expression of a target nucleic acid (RNA) sequence via binding of the compounds with the RNA sequence. The method comprises generating a library of virtual compounds *in silico* according to defined criteria, and evaluating *in silico* the binding of the virtual compounds with the RNA according to defined criteria. Also described are: (1) a method of defining a set of oligonucleotides (ONS) that modulate the expression of a RNA sequence via binding of the ONS with the RNA sequence comprising generating a library of virtual compounds *in silico* according to defined criteria, and evaluating the binding of the virtual ONS with the RNA according to defined criteria; and (2) a method of defining a set of compounds that modulate the expression of a RNA sequence via binding of the compounds with the RNA. The methods can be used for the generation and identification of synthetic compounds having defined physical, chemical or biocactive properties. Information gathered from assays of such compounds is used to identify nucleic acid sequences that are tractable to a variety of nucleotide sequence-based technologies, e.g., antisense drug discovery and target validation. AA240852 to AA2411220, and AAY52701 to AAY52706, represent sequences used in the

CC exemplification of the present invention.
CC
XX SQ Sequence 604 AA;

Query Match Similarity 100.0%; Score 294; DB 20; Length 604;
Best Local Similarity 100.0%; Pred. No. 2.7e-28; Mismatches 0; Indels 0; Gaps 0;
Matches 48; Conservative 0;

Qy 1 PEGIASGFYVGNSDVKCFCGGLRCMFGSDPPWQHAKWPRCE 48
Db 273 pegasagfyyvgnsdavkfcfcodgglcweesgdapwqhakwprce 320 .

RESULT 7

AAY33997 standard; Protein: 604 AA.
XX
AC XX
XX
DT 26-NOV-1999 (first entry)

DE Human cellular inhibitor of apoptosis-2 sequence.
KW cellular inhibitor of Apoptosis-2; antisense; diagnostic; therapeutic;
c-IAP-2; prophylaxis; infection; inflammation; tumor formation.
OS Homo sapiens.
XX
PN US5938771-A.
XX
PD 28-SEP-1999.
XX
PR 03-DEC-1998; 98US-0205144.
XX
PR 03-DEC-1998; 98US-0205144.
XX
PA (ISIS-) ISIS PHARM INC.
XX
PI Bennett CF, Cowser LM, Ackermann EJ;
DR N-PSDB; AA222096.
XX
WPI; 1999-561046/47.

PT Antisense compounds complementary to Cellular Inhibitor of Apoptosis-2
PT useful for e.g. diagnostics, therapeutics, and as research reagents.
XX
PS Example 13; Columns 45-50; 33pp; English.

CC The invention provides antisense compounds of 8-30 nucleotides that
CC inhibit the expression of human cellular inhibitor of apoptosis-2
(c-IAP-2). The antisense compounds may be used for diagnostics,
CC therapeutics (for modulating the expression of c-IAP-2), prophylaxis
CC (e.g. to prevent or delay infection, inflammation, or tumor formation),
CC as research reagents (e.g. to distinguish between members of a biological
CC pathway) and in kits. The present sequence represents the human cellular
CC inhibitor of apoptosis-2.
XX
SQ Sequence 604 AA;

Query Match Similarity 100.0%; Score 294; DB 20; Length 604;
Best Local Similarity 100.0%; Pred. No. 2.7e-28; Mismatches 0; Indels 0; Gaps 0;
Matches 48; Conservative 0;

Qy 1 PEGIASGFYVGNSDVKCFCGGLRCMFGSDPPWQHAKWPRCE 48
Db 273 pegasagfyyvgnsdavkfcfcodgglcweesgdapwqhakwprce 320 .

RESULT 8

AB50694 standard; Protein: 1141 AA.
ID AB50694

XX		DE	Human c- <i>AP1</i> repeat 3.
AC	AAB50694;	KX	
XX	19-MAR-2001 (first entry)	KW	TAP; inhibitor; apoptosis; RING finger domain; restinosis;
DT		XX	myocardial infarction; nephritis; HIV.
DE	Human AP12-MLT chimeric protein sequence.	OS	Homo sapiens.
XX		XX	
KW	Human; AP12-MLT chimera; apoptosis inhibitor 2; MLT; AP12;	PN	W09706182-A1.
KW	mucosa-associated lymphoid tissue	PD	20-FEB-1997.
KW	chromosome 11 region q21-22; 3; chromosome 18 region q21-1-22;	XX	06-AUG-1996;
KW	molecular characterisation; chromosome translocation; carcinogenesis;	PP	96WO-US12860.
XX	fusion protein; malignancy.	XX	08-DEC-1995;
OS		PR	95US-0569749.
OS	Chimeric - Homo sapiens.	PR	08-AUG-1995;
Synthetic.		XX	95US-0512946.
XX		PA	(TULA-) TULARIK INC.
PN	WO200073500-A1.	XX	
XX		PA	
PD	07-DEC-2000.	XX	
XX		PT	
PR	26-MAY-2000; 2000WO-EP04796.	XX	
XX		DR	WPI; 1997-154209/14.
PR	27-MAY-1999; 99EP0-0201683.	XX	
XX		PT	Nucleic acids encoding cellular inhibitor of apoptosis proteins -
PA	(VLA>) VLAMM INTERUNIVERSITAIR INST BIOTECHNOG.	PT	useful for apoptosis regulation in cells to reduce or increase
XX		XX	apoptosis and for pharmacological screening
PI	Baens M, Marynen P, Dierlam J;	FS	
XX		XX	
DR	WPI; 2001-061556/07.	CC	Claim 3; Page 25; 35pp; English.
DR	N-PSDB; AAC0972.	CC	The human cellular inhibitor of apoptosis proteins (c- <i>AP1</i> 1/2 -
XX		CC	repeats and/or a RING finger domain; in particular, at least two of
XX	determining if a tissue sample has a chromosome (11:18) translocation	CC	a first domain repeat (AAW1347 or AAW1350), a second domain repeat
PT	associated with malignancies by amplifying a nucleic acid sample using	CC	(AAW13549, AAW13550), and a third domain repeat (AAW13551 or AAW13552)
PT	primers complementary to chromosome 11 region q21-22.3 and chromosome	CC	and/or a RING finger domain (AAW13533 or AAW13554), or a consensus
PT	18 region q21.1-22.	CC	sequences derived from these human genes.
XX		CC	The nucleic acid is used for recombinant produ. of human cellular
PS	Claim 12; FIG 5; 47pp; English.	CC	inhibitor of apoptosis protein which modulates apoptosis
XX		CC	regulation. The nucleic acids are useful in therapies where
CC	comprises a chromosome (11:18) translocation associated with malignancies	CC	increased cell-specific apoptosis is desired, e.g. in restinosis,
CC	such as mucosa-associated lymphoid tissue lymphomas. The nucleic acid or	CC	inflammatory disease states, myocardial infarction, glomerular
CC	the antibody may be used as a probe for detection, for hybridisation to	CC	nephritis, transplant rejection and infectious diseases, e.g. HIV.
CC	southern blot, cell DNAs or for in situ hybridisation of cells, or for	CC	They can also be used in conditions requiring a reduction in
CC	determining the presence of complementary DNA. The present sequence	CC	apoptosis.
CC	represents the specifically claimed chimeric human apoptosis inhibitor 2	XX	
CC	(AP12)/MLT-lymphoma associated translocation (MLT) protein.	SO	Sequence 48 AA:
XX			
SQ	Sequence 1141 AA;		
Query Match Score 95.9%; DB 18; Length 48;			
Best Local Similarity 93.8%; Pred. No. 5.3e-26; 1; Indels 0; Gaps 0;			
Matches 45; Conservative 2; Mismatches 1;			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 273 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 10			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 48			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			
RESULT 9			
Matched保守性 0; 错配数 0; 缺失数 0; 插入数 0; 空缺数 0;			
OY 1 PEQLASAGFYYGVNSDDVKCFCGGGLRCKMESGDPWQHAKWPRCE 48			
Db 1 peqlasagfyygvnsddvkfcfcggglrcmesgdpwqhakwprce 320			

XX								
PD	14-NOV-1996.							
XX								
PF	11-MAY-1995;	95WO-US05922.						
XX								
PR	11-MAY-1995;	95WO-US05922.						
XX								
PA	(HUMA-)	HUMAN GENOME SCI INC.						
XX								
PT	He WM,	Hudson PL,	Rosen CA;					
XX								
DR	WPI; 1996-518608/51.							
DR	N-PSDB;	AAT43709.						
XX								
PT	Polyucleotide encoding human inhibitor of apoptosis gene 1 - useful							
PT	for treating degenerative diseases, as antiviral defence mechanism							
PT	and preventing cell death during trauma and strokes							
XX								
PS	Claim 1;	Page 40-41;	53pp;	English.				
XX								
CC	Human inhibitor of apoptosis 1 (hIAP-1) (AAW04583) is a protein							
CC	useful for treating degenerative diseases, rheumatoid arthritis,							
CC	septic shock, as an antiviral defence mechanism, and for							
CC	preventing cell death during strokes or trauma. Its amino acid							
CC	sequence was deduced from a cDNA clone (AAW43709) that can be obtai							
CC	from human Jurkat cell lines or human osteoclastoma stromal cell							
CC	lines. Recombinant hIAP-1 can be produced in prokaryotic or							
CC	eukaryotic host cells, or expressed in vivo. It can also be used							
CC	to screen for modulators of hIAP-1 activity.							
XX								
SQ	Sequence	438 AA;						
Query Match	95.9%	Score 282;	DB 17;	Length 438;				
Best Local Similarity	93.8%	Pred. No. 6e-27;	1;	Indels 0;	Gaps 0;			
Matches	45;	Conservative	2;	Mismatches 1;				
QY	1	PEQLASAGFYYVGNSDVYKFCFGDGRLRWESGGDPWPWHAKWFRC 48						
Db	107	peqlasagfyyvgnndvkfcfdggirclwesggdpwwehakwfrc 154						
RESULT 11								
AAW19746								
ID	AAW19746	standard;	Protein;	618 AA.				
AC	AAW19746;							
XX								
DT	16-SEP-1997	(first entry)						
XX								
DE	Human inhibitor of apoptosis protein homologue MIHB.							
KW	Inhibitor of apoptosis protein; IAP; mammalian IAP homologue; MIHB;							
KW	degenerative disease; infectious disease; autoimmune disease;							
KW	cancer; therapy; diagnosis.							
XX								
OS	Homo sapiens.							
XX								
PH	Key	Location/Qualifiers						
FT	Region	46..113						
FT		/label= BIR						
FT	Region	184..250						
FT		/label= BIR						
FT	Region	269..337						
FT		/label= BIR						
FT	Region	569..505						
FT		/label= RING_finger						
XX								
PN	W09723501-A1.							
XX								
PD	03-JUL-1997.							
XX								
PF	20-DEC-1996;	96WO-AU00827.						
XX								
Query Match	95.9%	Score 282;	DB 18;	Length 618;				
Best Local Similarity	93.8%	Pred. No. 8.6e-27;	1;	Indels 0;	Gaps 0;			
Matches	45;	Conservative	2;	Mismatches 1;				
QY	1	PEQLASAGFYYVGNSDVYKFCFGDGRLRWESGGDPWPWHAKWFRC 48						
Db	287	peqlasagfyyvgnndvkfcfdggirclwesggdpwwehakwfrc 334						
RESULT 12								
AAW1345								
ID	AAW1345	standard;	Protein;	618 AA.				
AC	AAW1345;							
XX								
DT	22-JUL-1997	(first entry)						
XX								
DE	Human c-IAP1.							
XX								
KW	IAP; inhibitor; apoptosis; RING finger domain; restinosis;							
KW	myocardial infarction; nephritis; HIV.							
OS	Homo sapiens.							
XX								
PN	W09706182-A1.							
XX								
PD	20-FEB-1997.							
XX								
PF	06-AUG-1996;	96WO-US12860.						
XX								
PR	08-DEC-1995;	95US-0562749.						
PR	08-AUG-1995;	95US-0512949.						
XX								
PA	(TULA-)	TULARIK INC.						
XX								
PT	Goeddel DV,	Rothe M;						
XX								
DR	WPI; 1997-154209/14.							
DR	N-PSDB;	AAT61590.						
XX								
PT	Nucleic acids encoding cellular inhibitor of apoptosis proteins -							
PT	useful for apoptosis regulation in cells to reduce or increase							

PT	apoptosis and for pharmacological screening	CC	pathway) and in kits. The present sequence represents the human cellular
PS	Disclosure; Page 18-20; 35pp; English.	XX	inhibitor of apoptosis-1.
XX	The human cellular inhibitor of apoptosis proteins (c-IAP1/2 -	CC	Sequence 618 AA;
CC	repeats and/or a RING finger domain; in particular, at least two of	CC	Query Match Similarity 95.9%; Score 282; DB 20; Length 618;
CC	a first domain repeat (AAW13547 or AAW13548), a second domain repeat	CC	Best Local Similarity 93.8%; Pred. No. 8; 5e-27; 1; Index 0; Gaps 0;
CC	(AAW13549 or AAW13550), and a third domain repeat (AAW13551 or AAW13552)	CC	Mismatches 45; Conservative 2; Mismatches 1; Index 0; Gaps 0;
CC	and/or a RING finger domain (AAW13553 or AAW13554), or a consensus	CC	Matches 45; Conservative 2; Mismatches 1; Index 0; Gaps 0;
CC	sequences derived from these human genes.	CC	Query 1 PEQQASAGFYYVGNSDDVKCFCGSGLARWEQGDPPWQHAKWPRCE 48
CC	The nucleic acid is used for recombinant prodn. of human cellular	CC	Db 287 peqasagfyyvgndvkcfccggircwesqgdppvhakwprce 334
CC	inhibitor of apoptosis protein which modulates apoptosis	CC	
CC	regulation. The nucleic acids are useful in therapies where	CC	
CC	increased cell-specific apoptosis is desired, e.g. in retinosis,	CC	
CC	inflammatory disease states, myocardial infarction, glomerular	CC	
CC	nephritis, transplant rejection and infectious diseases, e.g. HIV.	CC	
CC	They can also be used in conditions requiring a reduction in	CC	
CC	apoptosis.	CC	
SQ	Sequence 618 AA;		
RESULT 13			
Best Local Similarity 95.9%; Score 282; DB 18; Length 618;			
Matches 45; Conservative 2; Mismatches 1; Index 0; Gaps 0;			
Query 1 PEQQASAGFYYVGNSDDVKCFCGSGLARWEQGDPPWQHAKWPRCE 48			
Db 287 peqasagfyyvgndvkcfccggircwesqgdppvhakwprce 334			
RESULT 14			
AAW19583 ID AAW19583 standard; Protein: 618 AA.			
XX			
AC AAW19583;			
XX			
DT 02-SEP-1997 (first entry)			
DT			
DE Human apoptosis inhibitor HIAP-2.			
XX			
KW Apoptosis inhibitor; HIAP-2; HIV; AIDS; neurodegeneration;			
KW myelodysplastic syndrome; ischemia; myocardial infarction; stroke;			
KW reperfusion injury; toxin-induced liver disease; gene therapy;			
KW diagnosis.			
XX			
OS Homo sapiens.			
XX			
PH Key	Location/Qualifiers		
PT Domain	46..113 /label=BIR-1		
PT	184..220 /label=BIR-2		
PT Domain	269..336 /label=BIR-3		
PT	560..605 /label=Ring_zinc_finger		
PT Domain			
PN WO9706255-A2.			
XX			
PD 20-FEB-1997.			
XX			
PF 05-AUG-1996; 96WO-1B01022.			
XX			
PR 22-DEC-1995; 95US-0576956.			
XX			
PR 04-AUG-1995; 95US-0511485.			
XX			
PA (UWO ⁺) UONV OTTAWA.			
XX			
PI Baird S, Korneluk RG, Liston P, Mackenzie AE;			
XX			
DR WPI.1997-15-262/14.			
DR N-PSDB; AAT70838.			
XX			
PT Nucleic acid encoding an inhibitor of apoptosis polypeptide - used			
PT to inhibit apoptosis in e.g. HIV or AIDS patients, and for detection			
PT of susceptibility to apoptotic disease			
XX			
PS Claim 27; Page 75-77; 219pp; English.			
XX			
CC Human XIAP, HIAP-1 and HIAP-2, and murine XIAP, M-HIAP-1 and			
CC M-HIAP-2 (AAW19581-6-6) are a new class of mammalian proteins that			
CC are inhibitors of apoptosis (IAP) and which are characterised by			
CC the presence of a ring zinc finger domain (see also AAW19587) and at			
CC least one BIR (baculovirus IAP repeat) domain (see also AAW19588).			
CC The IAP amino acid sequences were deduced from cDNA clones (AAW70837			
CC and AAW70838) from a human liver library. IAP polypeptides can be			
CC expressed in host cells (in vitro or in vivo) and used in methods for			
CC treating diseases and disorders involving apoptosis, esp. in a			

Example 13: Columns 41-46; 32pp; English.

The invention provides antisense compounds of 8-30 nucleotides that inhibit the expression of human Cellular Inhibitor of Apoptosis-1 (c-IAP-1). The antisense compounds may be used for diagnostics, therapeutics (for modulating the expression of c-IAP-1), prophylaxis

(e.g. to prevent or delay infection, inflammation, or tumor formation), as research reagents (e.g. to distinguish between members of a biological

CC human diagnosed as HIV-positive or as having AIDS, a neurodegenerative disease, a myelodysplastic syndrome or an ischemic injury, selected from myocardial infarction, stroke, reperfusion injury, or a toxin-induced liver disease.

CC Sequence 618 AA:

SQ Query Match 91.8%; Score 270; DB 19; Length 618; Best Local Similarity 91.7%; Pred. No. 2, 6e-25; Matches 44; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 1 PEOIASAGFYVGNSDVKFCFCDGGLRKGESGDPWYHAKWFPRCE 48
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 287 peqlasagfyvgrndvckfcgddgglrcwesgdppwakwfprce 334

RESULT 15

AAW69296 15

ID AAW69295 standard; Protein: 618 AA.

XX

AC AAW69296;

XX DT 13-NOV-1998 (first entry)

XX DE Human HIAP-2 protein.

XX KW Inhibitor of apoptosis protein; apoptosis enhancer; NAIp polypeptide; proliferative disease; IAP; therapy; cancer; human; HIAP-2 protein.

XX OS Homo sapiens.

XX PN W09815693-A2.

XX PD 20-AUG-1998.

XX PF 13-FEB-1998;

XX PR 13-FEB-1997; 970US-0800929.

XX PA (UYOP-) UNIV OTTAWA.

XX PI Baird S, Korneluk R, Liston P, Mackenzie AE, Pratt C;

PI Tsang B;

XX DR WPI: 1998-467164/40.

XX DR N-PSDB; AAV55040.

PT Inducing apoptosis in proliferative mammalian cells with inhibitor of IAP or NAIP polypeptide - also methods for prognosis based on presence of IAP and NAIP specifically applied to cancers involving PT p53 mutations

PT p53 mutations

PS Disclosure: FIG 3: 14TPP: English.

XX

CC This sequence is the human HIAP-2 protein, which is a inhibitor of apoptosis protein (IAP), and can be used in the method of the invention. The method is for enhancing apoptosis in cells from a mammal with proliferative disease by treatment with a compound that inhibits biological activity of an IAP or NAIP polypeptide. The inhibitory compounds are used to treat proliferative diseases, specially cancers of ovary, breast, pancreas, lymph nodes, skin, blood, lung, brain, kidney, liver, nasopharynx, thyroid, central nervous system, prostate, colon, rectum, cervix or endometrium, particularly to increase their sensitivity to chemotherapeutic agents. High levels of the IAP or NAIP proteins are detected in many cancers and are associated with poor prognosis, resistance to chemotherapeutic agents and mutations in p53 (it is suggested that 14TPP, p53 suppresses transcription of the IAP or NAIP genes). Transgenic animals are used for testing the effects of antisense CC oligonucleotides and for screening for the inhibitors.

CC Sequence 618 AA:

SQ Sequence 618 AA:

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